

Applicant : Fan, Jian-Qiang
Serial No. : 10/781,356 Examiner : Schnizer, Richard A.
Filed : February 17, 2004 Group Art Unit : 1635
Customer No. 21003 Confirmation No. 9219
For : COMBINATION THERAPY FOR TREATING PROTEIN
DEFICIENCIES

SUPPLEMENTAL INFORMATION DISCLOSURE STATEMENT

Commissioner for Patents
P.O. Box 1450
Alexandria, VA 22313-1450

Sir:

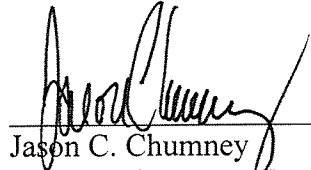
Pursuant to 37 C.F.R. §§ 1.56 and 1.97, applicant brings to the attention of the Examiner the document listed on the attached Form PTO 1449 and respectfully request that the listed document be considered by the Examiner and made of record in the above-captioned application.

This submission does not represent that a search has been made or that no better art exists and does not constitute an admission that the listed document is material or constitutes "prior art." If the Examiner applies the document as prior art against any claim in the application and applicant determines that the cited document do not constitute "prior art" under United States law, applicant reserves the right to present to the Office the relevant facts and law regarding the appropriate status of the document.

Applicant further reserves the right to take appropriate action to establish the patentability of the disclosed invention over the listed document, should the document be applied against the claims of the present application.

Applicant believes no fee is due. If any additional fee is due, or if any overpayment has been made, the Commissioner is authorized to charge any such fee or credit any overpayment, to our Deposit Account No. 02-4377.

Respectfully submitted,



Jason C. Chumney
Patent Office Reg. No. 54,781

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Enclosure

Substitute for form 1449/PTO INFORMATION DISCLOSURE STATEMENT BY APPLICANT (Use as many sheets as necessary)	Application Number	10/781,356
	Filing Date	February 17, 2004
	Inventor(s)	Fan
	Art Unit	1635
	Examiner Name	Schnizer
	Attorney Docket Number	070165.0633

Exam Initial	No.	OTHER DOCUMENTS (including Author, Title, Date, Pertinent Pages, Etc.)
	1.	Handa et al., 2000, "A case of symptomatic heterozygous female Fabry's disease without detectable mutation in the alpha-galactosidase gene," Dermatology 200:262-265

Examiner Signature		Date Considered	
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NY02:596630.1

* Examiner: Initial citation considered, whether or not citation is in conformance with MPEP 609; Draw line through citation if not in conformance and not considered. Include copy of this form with next communication to applicant.